WAL ZYR D- cetirizine hydrochloride, pseudoephedrine hydrochloride tablet, film coated, extended release Walgreen Company

Walgreen Co. Wal-Zyr [™] D Drug Facts

Active ingredients (in each extended release tablet)

Cetirizine HCl 5 mg

Pseudoephedrine HCl 120 mg

Purpose

Antihistamine Nasal decongestant

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat
- nasal congestion
- reduces swelling of nasal passages
- temporarily relieves sinus congestion and pressure
- temporarily restores freer breathing through the nose

Warnings

Do not use

- if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing hydroxyzine.
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- heart disease
- thyroid disease
- diabetes
- glaucoma
- high blood pressure
- trouble urinating due to an enlarged prostate gland

• liver or kidney disease. Your doctor should determine if you need a different dose.

Ask a doctor or pharmacist before use if you are

taking tranquilizers or sedatives.

When using this product

- do not use more than directed
- drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

- an allergic reaction to this product occurs. Seek medical help right away.
- you get nervous, dizzy, or sleepless
- symptoms do not improve within 7 days or are accompanied by fever

If pregnant or breast-feeding:

- if breast-feeding: not recommended
- if pregnant: ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

• do not break or chew tablet; swallow tablet whole

adults and children 12 years and over	take 1 tablet every 12 hours; do not take more than 2 tablets in 24 hours.
adults 65 years and over	ask a doctor
children under 12 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- store between 20° to 25°C (68° to 77°F)
- do not use if blister unit is broken or torn

- see side panel for lot number and expiration date
- meets USP Dissolution Test 2

Inactive ingredients

colloidal silicon dioxide, hypromellose, lactose monohydrate, low-substituted hydroxypropyl cellulose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, talc, titanium dioxide

Questions or comments?

1 - 800 - 719 - 9260

Package/Label Principal Display Panel

Wal-Zyr™D

Compare to Zyrtec-D[®]

active ingredients

12 HOUR ALLERGY & CONGESTION

CETIRIZINE HYDROCHLORIDE &

PSEUDOEPHEDRINE HYDROCHLORIDE

EXTENDED-RELEASE TABLETS 5 mg / 120 mg

ANTIHISTAMINE / NASAL DECONGESTANT

12 HOUR EXTENDED-RELEASE TABLETS

INDOOR & OUTDOOR ALLERGIES

12-hour relief of sneezing; runny nose; itchy, throat or nose; itchy, watery eyes; sinus pressure & nasal congestion

12 EXTENDED-RELEASE TABLETS

12 HOUR

ACTUAL SIZE

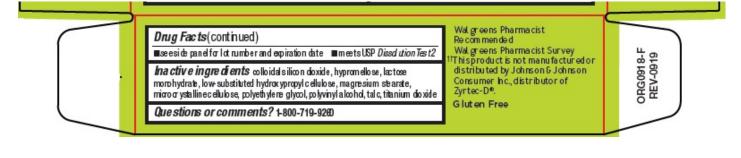


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CODE ARE/

Drug Facts	Drug Facts (continued)			
Active ingredients Purpose (in each extended release tablet)	When using this product I do not use more than directed			
Cetirizine HC15 mgAnîhis tamine Pseudcephedrine HC1120 mgNasal de corgestant				
Uses Hemporarily relieves these symptoms due to hay feveror other upper respiratory allergies Trunny nose sneezing tichy, watery eyes itchingof the rose or throat to as all congestion reduces swelling of nasal pass ages temporarily relieves sinus congestion and pressure				
■ temporarily restores freer breathing through the nose				
<i>Warnings</i> Donotuse ■ ifyouhave ever hadanallergic reaction to thisproductor any				
ofits ingredients or to a nantihistamin econtaining hydroxyzine. ■ if you are now taking a prescription monoamine oxidas e	Directions do not break or chew tablet, swalow tablet whole			
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	adults 65 years and over askadoctor			
before takingthis product.	children un der ask adoctor 12yearsofage			
Ask a doctor before use if you have ■ heartdise ase ■ thyroid dise ase ■ diabetes ■ glaucom a ■ high blood pressure	consumers with liver askadoctor or kidney disease			
 trouble urina ing due to an enlarged prostate gland liver or kidney dise ase. Your doctor should determine if you need a different dos e. 	Ot he rinform ation ■storebetween 20° to 25°C (68° to 77°F) ■donotuse if blister unit is broken ortorn ▼			

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WAL ZYR D

cetirizine hydrochloride, pseudoephedrine hydrochloride tablet, film coated, extended release

Product Informatio	n							
Product Type		HUMAN OTC DRI	UG	Item Code (Source)		NDC:0363-	NDC:0363-1147	
Route of Administratio	n	ORAL						
Active Ingredient/A	Active Moie	ty						
	Ingre	dient Name			Basis of St	rength	Strength	
CETIRIZINE HYDRO CHLO RIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:YO7261ME24)				CETIRIZINE HYDROCHLORIDE		5 mg		
PSEUDO EPHEDRINE HY - UNII:7CUC9DDI9F)	DROCHLORI	DE (UNII: 6 V9 V2F	RYJ8N) (PSEU	DOEPHEDRINE	PSEUDOEPHEDRINE HYDROCHLORIDE		120 mg	
Inactive Ingredients								
	Ingredient Name					Strength		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)								
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)								
LACTOSE MONOHYDR	ATE (UNII: EW	Q57Q8I5X)						
LOW-SUBSTITUTED HY	DRO XYPRO I	PYL CELLULOS	E, UNSPECIFI	ED (UNII: 2165F	E0K14)			
MAGNESIUM STEARAT	MAGNESIUM STEARATE (UNII: 70097M6I30)							
MICRO CRYSTALLINE C	MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)							
POLYETHYLENE GLYC	OL, UNSPECI	FIED (UNII: 3WJQ	0SDW1A)					
POLYVINYL ALCOHOL	., UNSPECIFIE	2 D (UNII: 532B59J	990)					
TALC (UNII: 7SEV7J4R1U	·							
TITANIUM DIO XIDE (UN	NII: 15FIX9V2JF	2)						
Product Characteri	istics							
Color	WHITE		Score		10 score			
Shape	ROUND)	Size		0 mm			
Flavor			Imprint Code L147		147	7		
Contains								
Packaging								
# Item Code Package Description Man			keting Start Date	Marketin	g End Date			
1 NDC:0363-1147-53 12 in 1 CARTON			03/26	/2020				

1	1 in 1 BLISTER PACK; Type 0: Not a Combination Product					
2 NDC:0363-1147-62	24 in 1 CARTON	03/26/2020				
2	1 in 1 BLISTER PACK; Type 0: Not a Combination Product					
Marketing Information						
Marketing Categor	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
ANDA	ANDA210719	03/26/2020				

Labeler - Walgreen Company (008965063)

Revised: 3/2020

Walgreen Company